

(19)



Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 0 926 515 A1

(12)

EUROPEAN PATENT APPLICATION

published in accordance with Art. 158(3) EPC

(43) Date of publication:

30.06.1999 Bulletin 1999/26

(51) Int. Cl.⁶: **G02B 5/30, C08J 5/18**

(21) Application number: **98907265.7**

(86) International application number:

PCT/JP98/01119

(22) Date of filing: **17.03.1998**

(87) International publication number:

WO 98/41887 (24.09.1998 Gazette 1998/38)

(84) Designated Contracting States:

DE FR GB IT

(72) Inventor: **ISOZAKI, Takanori**

Kurashiki-shi Okayama 710-8622 (JP)

(30) Priority: **19.03.1997 JP 6584297**

(74) Representative:

Nachshen, Neil Jacob et al

D Young & Co

21 New Fetter Lane

London EC4A 1DA (GB)

(71) Applicant: **Kuraray Co., Ltd.**

Kurashiki-City, Okayama 710-8622 (JP)

(54) **POLARIZING FILM**

(57) The present invention provides a polarizing film having a dichroic ratio of at least 20 and comprising a polyvinyl alcohol derivative with a polyvinylene structure as obtained from a polyvinyl alcohol material having a degree of polymerization of at least 2000.

The polarizing film of the present invention has excellent polarizing properties in the range of the visible ray spectrum, and exhibits excellent durability when left in high-temperature and high-humidity surroundings for a long period of time. The polarizing film of the present invention is useful for a polarizing sheet laminated with a protective film for on-vehicle LCD that is placed in heavy temperature and humidity fluctuations, such as LCD navigation systems, LCD TV, etc.

EP 0 926 515 A1

Description

Technical Field

[0001] The present invention relates to a polarizing film comprising a polyvinyl alcohol derivative with a polyvinylene structure.

Background Arts

[0002] A Polarizing sheet having a light-transmitting and shielding function laminated with protective films is a basic constituent element for a liquid crystal display (LCD), along with a liquid crystal having a light-switching function. The applications of the LCD include small-sized instruments such as pocket calculators, watches, etc. in the early years, as well as lap-top personal computers, word processors, liquid-crystal color projectors, on-vehicle navigation systems, liquid-crystal TV, indoor and outdoor measuring instruments, etc. The LCD has been used in heavy temperature and humidity fluctuations of from low temperature to high temperature and from low humidity to high humidity. Accordingly, there has been desired a polarizing sheet which has excellent durability such as wet heat resistance, etc. and excellent polarizing properties.

[0003] A polarizing sheet has a structure that both sides of a polarizing film as obtained from a polyvinyl alcohol (hereinafter referred to as PVA) film are laminated with supporting films of cellulose triacetate or the like. As a polarizing film, there have been known an iodine-containing polarizing film as prepared by dyeing a PVA film with iodine, a dye-containing polarizing film as prepared by dyeing a PVA film with a dichroic dye, and a polarizing film of a PVA derivative with a polyvinylene structure as prepared by dehydrating a PVA film (U.S. Patent No. 2,173,304).

[0004] The iodine-containing polarizing film and the dye-containing polarizing film have good polarizing properties, while they have a problem that their durability is poor. The polarizing film of a polyvinyl alcohol derivative with a polyvinylene structure has excellent durability, while it has a problem that its polarizing property is poor.

Disclosure of the Invention

[0005] The object of the present invention is to provide a polarizing film which has excellent durability and excellent polarizing properties.

[0006] Having intensively studied to solve the above object, the present inventor has found out a polarizing film having a dichroic ratio of at least 20 and comprising a polyvinyl alcohol derivative with a polyvinylene structure as obtained from a polyvinyl alcohol material having a degree of polymerization of at least 2000, and has completed the present invention.

Best Modes of Carrying out the Invention

[0007] The present invention is described in more detail hereinafter.

[0008] The degree of polymerization of the PVA material to be used for producing the polarizing film of the present invention is at least 2000, but preferably at least 2800, more preferably at least 3000, even more preferably at least 3500. The higher the degree of polymerization, the better the durability and the polarizing properties. In view of the film-forming properties, the upper limit of the degree of polymerization is preferably 30000. The degree of polymerization of the PVA is measured according to JIS-K-6726.

[0009] In view of the polarizing properties and the durability, the degree of hydrolysis of the PVA is preferably at least 90 mol%, more preferably at least 98 mol%, even more preferably at least 99 mol%, still more preferably at least 99.5 mol%.

[0010] The dichroic ratio of the polarizing film is at least 20, preferably at least 25, more preferably at least 30. To increase its dichroic ratio, it is desirable that a PVA film having a high degree of polymerization and containing a dehydrating promoter is stretched under dry-heat at a high stretch ratio in an oxygen-poor atmosphere.

[0011] The wavelength of maximal absorption in the range of the visible ray absorption spectrum (wavelength: 380 to 780 nm) of the polarizing film is preferably at least 500 nm, more preferably at least 520 nm, even more preferably at least 540 nm. The upper limit of the wavelength of maximal absorption is preferably 650 nm, more preferably 620 nm. As the degree of polymerization of the PVA to be used as a material becomes higher, the wavelength of maximal absorption tends to shift to a higher wavelength side.

[0012] In the case where the wavelength of maximal absorption is in the above-described range, it is estimated that the number of vinylene units of conjugated double bonds which constitute the polyvinylene structure be approximately from 15 to 30.

[0013] As a method for producing PVA, for example, there are a method of hydrolyzing a polyvinyl ester polymer as obtained through radical polymerization of a vinyl ester monomer such as vinyl acetate and the like, with an alkali catalyst or an acid catalyst.

[0014] As a method for polymerizing a vinyl ester monomer, for example, there are bulk polymerization, solution polymerization, suspension polymerization, emulsion polymerization and the like. Bulk polymerization or emulsion polymerization is preferable to obtain PVA having a degree of polymerization of at least 4000. As a polymerization catalyst, for example, there are azo catalyst, peroxide catalyst, redox catalyst, etc.

[0015] As a vinyl ester monomer, for example, there are vinyl formate, vinyl acetate, vinyl propionate, vinyl valerate, vinyl caprate, vinyl laurate, vinyl stearate, vinyl benzoate, vinyl pivalate, vinyl trifluoroacetate, vinyl ver-

satate, etc. Of those, vinyl acetate is preferable.

[0016] The vinyl ester monomer may be copolymerized with a copolymerizable comonomer.

[0017] As a comonomer, for example, there are olefins such as ethylene, propylene, 1-butene, isobutene, etc.; acrylic acid and its salts; acrylates such as methyl acrylate, ethyl acrylate, n-propyl acrylate, i-propyl acrylate, n-butyl acrylate, i-butyl acrylate, t-butyl acrylate, 2-ethylhexyl acrylate, dodecyl acrylate, octadecyl acrylate, etc.; methacrylic acid and its salts; methacrylates such as methyl methacrylate, ethyl methacrylate, n-propyl methacrylate, i-propyl methacrylate, n-butyl methacrylate, i-butyl methacrylate, t-butyl methacrylate, 2-ethylhexyl methacrylate, dodecyl methacrylate, octadecyl methacrylate, etc.; acrylamide derivatives such as acrylamide, N-methylacrylamide, N-ethylacrylamide, N,N-dimethylacrylamide, diacetoneacrylamide, acrylamidopropanesulfonic acid and its salts, acrylamidopropylidimethylamine and its salts and quaternary salts, N-methylolacrylamide and its derivatives, etc.; methacrylamide derivatives such as methacrylamide, N-methylmethacrylamide, N-ethylmethacrylamide, methacrylamidopropanesulfonic acid and its salts, methacrylamidopropylidimethylamine and its salts and quaternary salt, N-methylolmethacrylamide and its derivatives, etc.; vinyl ethers such as methyl vinyl ether, ethyl vinyl ether, n-propyl vinyl ether, i-propyl vinyl ether, n-butyl vinyl ether, i-butyl vinyl ether, t-butyl vinyl ether, dodecyl vinyl ether, stearyl vinyl ether, etc.; nitriles such as acrylonitrile, methacrylonitrile, etc.; vinyl halides such as vinyl chloride, vinylidene chloride, vinyl fluoride, vinylidene fluoride, etc.; allyl compounds such as allyl acetate, allyl chloride, etc.; maleic acid and its salts and esters; vinylsilyl compounds such as vinyltrimethoxysilane, etc.; isopropenyl acetate, etc.

[0018] As another method for producing PVA, there may be a method of hydrolyzing a polyvinyl ether polymer from t-butyl vinyl ether, benzyl vinyl ether, trimethylsilyl vinyl ether and the like.

[0019] A PVA film as prepared by film-forming a PVA having a degree of polymerization of at least 2000 is used for producing the polarizing film of the present invention. As a method of film-forming a PVA film, for example, there is a casting method or a die-casting method that comprises casting a PVA solution onto a resin film, a drying drum or a drying belt. As a solvent for the PVA, for example, water, an organic solvent, and a mixed solvent of water and an organic solvent. The organic solvent includes dimethylsulfoxide, phenol, methanol, ethanol, etc. The PVA solution may optionally contain any of plasticizers, surfactants, dichroic dyes, inorganic salts, etc. The PVA film may be heat-treated, if desired. The thickness of the PVA film is preferably 5 to 500 μm , more preferably 30 to 200 μm .

[0020] The process for producing the polarizing film comprises a treatment of pre-swelling a PVA film, treatment of dehydrating so as to form a polyvinylene structure in the PVA film, a treatment of monoaxially

stretching (in many cases, the monoaxial stretching and the dehydration are often conducted at the same time), a treatment of fixing with a boron compound or the like, and a treatment of drying. If desired, a heat treatment may be conducted. There is no limitation on the order and the number of each treatment.

[0021] As the treatment of dehydrating so as to form a polyvinylene structure in the PVA film, for example, there are a method of stretching a PVA film that contains a dehydrating promoter, and a method of heating a stretched PVA film with a dehydrating promoter. As a method of adding a dehydrating promoter into the PVA film, there are a method of dipping the PVA film in an aqueous solution containing a dehydrating promoter, a method of placing the PVA film in an atmosphere of a gaseous dehydrating promoter, and a method of film-forming the PVA solution containing a dehydrating promoter. As a dehydrating promoter, there are protonic acids such as hydrochloric acid, sulfuric acid, etc.; and stannous halides such as stannous chloride, stannous bromide, etc. The amount of the dehydrating promoter is preferably 0.001 to 10 % by weight based on the PVA. In the case where the amount of the dehydrating promoter is less than the above-described range, the polarizing performance hardly appears. On the other hand, in the case where it is more than the above-described range, the efficient dehydration becomes difficult. The dehydrating promoter may be added to the PVA film during the film-forming step, before the monoaxial stretching step, or before the dehydrating step. When the dehydrating promoter is added to the film-forming stock solution for the PVA film, the care should be paid because too high drying temperature during the film-forming causes the possibility of the thermal oxidation of PVA.

[0022] As a method of monoaxial stretching, for example, there are dry-heat stretching, wet stretching, stretching in warm water, stretching in air with the conditions of water absorption.

[0023] The stretching temperature for dry-heat stretching is preferably at least 100°C, more preferably at least 120°C, even more preferably at least 140°C. The upper limit of the stretching temperature is preferably 250°C, more preferably 230°C, even more preferably 220°C. If desired, the stretching temperature may be changed, depending on the stretched film condition during stretching. The dry-heat stretching is preferably conducted in an oxygen-poor atmosphere such as a nitrogen atmosphere or in vacuum, because there is a possibility of discoloration due to the oxidation of PVA. The PVA film is converted into a colored film having polarizing properties during the dry-heat stretching.

[0024] The stretching temperature for wet stretching is preferably at least 20°C, more preferably at least 40°C, even more preferably at least 50°C. The upper limit of the stretching temperature is preferably 90°C, more preferably 80°C. A heat treatment is required because intended polyvinylene structure could not be formed

according to only wet stretching.

[0025] In the case where the stretching temperature is too low, the stretching with a high stretching ratio becomes difficult, resulting in that the polarizing properties could hardly be improved. On the other hand, in the case where the stretching temperature is too high, the film tends to be cut off during stretching. When the stretching is intermittently conducted two times or more, different stretching methods may be employed.

[0026] The stretching ratio is preferably at least 4-fold, more preferably at least 5-fold. With the increase in the stretching ratio, the wavelength of maximal absorption shifts to a higher wavelength side, and the optical performance is improved. The upper limit of the stretching ratio is preferably 10-fold, more preferably 9-fold, in view of the possibility of uniform stretching.

[0027] In the case where the dehydration (the formation of polyvinylene structure) by stretching is not enough, a dehydrating promoter is further added, and the film is heated to be dehydrated. The temperature for the dehydration is preferably 90 to 180°C, more preferably 130 to 170°C.

[0028] The amount of water to be removed from PVA is preferably 1 to 5 % by weight based on the PVA film, where the film has a thickness of 30 to 100 µm. In the case where the amount of removed water from PVA is too small, the absorption in visible ray decreases, and enough polarizing performance does not appear. On the other hand, in the case where the amount of removed water is too much, the absorption in visible ray increases too much, causing the difficulty in obtaining a polarizing film with a suitable transparency, and furthermore intermolecular crosslinking occurs, reducing the number of conjugated double bonds or making the film brittle.

[0029] In the case where the dehydrating promoter remains in the polarizing film, there is a possibility that the polarizing properties change while left at a high temperature. Therefore, it is desirable to remove the dehydrating promoter by dipping in water, weak-alkaline aqueous solution, water-containing methanol, or aqueous solution of an inorganic substance such as saline or the like.

[0030] After the formation of a polyvinylene structure in PVA film, it is preferably dipped in a bath for fixing treatment containing 1 to 6 % by weight of a boron compound (e.g., boric acid). The fixing treatment may be conducted according to the required waterproof level. The temperature in the bath for fixing treatment is preferably 20 to 70°C. The drying temperature after taking out of the fixing bath is preferably 30 to 80°C. A heat treatment may be conducted at 50 to 150°C after drying.

[0031] The thickness of the polarizing film of the present invention is preferably 5 to 200 µm, more preferably 10 to 100 µm.

[0032] In order to obtain the furthermore enough resistance to water, the polarizing film of the present

invention may be laminated, either on its both sides or on its one side, with a protective film which is transparent and has a mechanical strength, thereby giving a polarizing sheet. As a protective film, there are used cellulose acetate film, acrylic film, polyester film, etc.

[0033] The present invention is described concretely hereinunder with the following Examples, which, however, are not intended to restrict the scope of the present invention. Unless otherwise specifically indicated, "%" and "part" in Examples are "% by weight" and "part by weight".

[0034] The polyvinylene structure in a polarizing film was identified by measuring the wavelength of maximal absorption in the range of the visible ray absorption spectrum (wavelength: 380 to 780 nm) through UV and visible ray spectrophotometer.

[0035] The transmittance, the degree of polarization and the dichroic ratio of a polarizing film were measured and calculated according to the Standard of the Electronic Industry Association of Japan (EIAJ) LD-201-1983, using a spectrophotometer with a C light source in the standard colorimetric system.

[0036] A Polarizing film is generally used as being laminated with a protective film, however in the following Examples a polarizing film alone without a protective film was tested to measure its properties.

Example 1:

[0037] A film of PVA (degree of polymerization 4000, degree of hydrolysis 99.9 mol%) having a thickness of 100 µm was dipped in an aqueous solution of sulfuric acid (0.01 N) at 20°C for 10 minutes. This was dried at 20°C for 30 minutes. This was stretched 6.8-fold in the monoaxial direction, in nitrogen at 190°C. Next, this was dipped in an aqueous solution of 4 % boric acid for 10 minutes, whereby sulfuric acid was removed and, at the same time, the stretched film was fixed. Finally, this was dried in blowing warm air at 40°C. Thus was obtained a polarizing film having a polyvinylene structure therein and having a thickness of 20 µm. The results of evaluation on the polarizing film are shown in Table 1.

Example 2:

[0038] A film of PVA (degree of polymerization 9000, degree of hydrolysis 99.6 mol%) having a thickness of 75 µm was dipped in an aqueous solution of sulfuric acid (0.01 N) at 20°C for 10 minutes. This was dried at 20°C for 30 minutes. This was stretched 6.5-fold in the monoaxial direction, in nitrogen at 220°C. Next, this was rinsed by distilled water. Finally, this was dried in blowing warm air at 40°C. Thus was obtained a polarizing film having a polyvinylene structure therein and having a thickness of 18 µm. The results of evaluation on the polarizing film are shown in Table 1.

Example 3:

[0039] A film of PVA (degree of polymerization 2600, degree of hydrolysis 99.7 mol%) having a thickness of 75 μm was dipped in an aqueous solution of sulfuric acid (0.1 N) at 20°C for 10 minutes. This was dried at 20°C for 30 minutes. This was stretched 5.2-fold in the monoaxial direction, in nitrogen at 160°C. Next, this was rinsed by distilled water. Finally, this was dried in blowing warm air at 40°C. Thus was obtained a polarizing film having a polyvinylene structure therein and having a thickness of 24 μm . The results of evaluation on the polarizing film are shown in Table 1

Comparative Example 1:

[0040] A film of PVA (degree of polymerization 1700, degree of hydrolysis 99.9 mol%) having a thickness of 100 μm was dipped in an aqueous solution of sulfuric acid (0.5 N) at 20°C for 10 minutes. This was dried at 20°C for 30 minutes. This was stretched 4.3-fold in the monoaxial direction, in air at 130°C. Next, this was rinsed by distilled water. Finally, this was dried in blowing warm air at 40°C. Thus was obtained a polarizing film having a polyvinylene structure therein and having a thickness of 51 μm . The results of evaluation on the polarizing film are shown in Table 1.

Comparative Example 2:

[0041] A film of PVA (degree of polymerization 4000, degree of hydrolysis 99.7 mol%) having a thickness of 75 μm was dipped in an aqueous dyeing solution (iodine 0.2 g/liter, potassium iodide 20 g/liter, boric acid 40 g/liter) at 30°C for 3 minutes. Next, this was stretched 4-fold in the monoaxial direction, in an aqueous solution of boric acid (40 g/liter) at 50°C, and then dipped in an aqueous processing solution (potassium iodide 20 g/liter, boric acid 40 g/liter, zinc chloride 10 g/liter) at 30°C for 4 minutes. Finally, this was dried in blowing warm air at 40°C. Thus was obtained an iodine-containing polarizing film (with no polyvinylene structure) having a thickness of 16 μm . The results of evaluation on the polarizing film are shown in Table 1.

Table 1

	Optical Properties before shelf test			Optical Properties after shelf test (shelf test Condition: 80°C, 90 % RH, 1000 hours)		
	Transmittance (%)	Degree of Polarization (%)	Dichroic Ratio	Wavelength of Maximal Absorption (nm)	Transmittance (%)	Degree of Polarization (%)
Example 1	42.9	98.8	33	580	43.7	95.1
Example 2	42.5	99.2	34	590	43.0	97.0
Example 3	36.4	99.6	20	560	37.8	96.0
Comparative Example 1	40.0	77.2	7	490	42.9	58.1
Comparative Example 2	43.4	99.1	38	—	90.7	0.0
						1

Industrial Applicability

[0042] The polarizing film of the present invention has excellent polarizing properties in the range of the visible ray spectrum, and exhibits excellent durability when left in high-temperature and high-humidity surroundings for a long period of time. The polarizing film of the present invention is useful for a polarizing sheet laminated with a protective film for on-vehicle LCD that is placed in heavy temperature and humidity fluctuations, such as LCD navigation systems, LCD TV, etc.

Claims

1. A polarizing film having a dichroic ratio of at least 20 and comprising a polyvinyl alcohol derivative with a polyvinylene structure as obtained from a polyvinyl alcohol material having a degree of polymerization of at least 2000.
2. The polarizing film as claimed in claim 1, which has a wavelength of maximal absorption of at least 500 nm in the range of the visible ray absorption spectrum.
3. The polarizing film as claimed in claim 1 or 2, wherein the polyvinyl alcohol has a degree of hydrolysis of at least 90 mol%.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP98/01119

A. CLASSIFICATION OF SUBJECT MATTER Int.Cl ⁶ G02B5/30, C08J5/18		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) Int.Cl ⁶ G02B5/30, C08J5/00-5/02, C08J5/12-5/22		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Jitsuyo Shinan Koho 1926-1997 Toroku Jitsuyo Shinan Koho 1994-1997 Kokai Jitsuyo Shinan Koho 1971-1997		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US, 2445555, A (Polaroid Corporation), July 20, 1948 (20. 07. 48) (Family: none)	1-3
A	US, 3914017, A (Polaroid Corporation), October 21, 1975 (21. 10. 75) (Family: none)	1-3
A	US, 4818624, A (Polaroid Corporation), April 4, 1989 (04. 04. 89) (Family: none)	1-3
A	JP, 2-256002, A (Polaroid Corp.), October 16, 1990 (16. 10. 90) & US, 4895769, A & EP, 357946, A & DE, 68917836, E	1-3
A	JP, 2-282201, A (Bayer AG.), November 19, 1990 (19. 11. 90) & US, 5051286, A & EP, 374627, A & DE, 3905027, A	1-3
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.		
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "I" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family		
Date of the actual completion of the international search June 11, 1998 (11. 06. 98)		Date of mailing of the international search report June 23, 1998 (23. 06. 98)
Name and mailing address of the ISA/ Japanese Patent Office		Authorized officer
Facsimile No.		Telephone No.

Form PCT/ISA/210 (second sheet) (July 1992)

INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP98/01119

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	JP, 1-105204, A (Unitika Ltd.), April 21, 1989 (21. 04. 89) & US, 5071906, A & EP, 297927, A & DE, 3889256, G	1, 3
P	JP, 9-178944, A (Bayer AG.), July 11, 1997 (11. 07. 97) & DE, 19650198, A1	1-3

Form PCT/JP98/01119 (continuation of second sheet) (July 1992)